



# California Medical Device Recall Information



## Recall Name

### Vascular Solutions Recalls All Unexpired Lots of Twin-Pass Dual Access Catheters Due to Potential for Excess Manufacturing Material

Recall Date	Product Description	Recalling Firm	Recall Reason
9/16/16	Twin-Pass Dual Access Catheters  Model Numbers: <ul style="list-style-type: none"><li>• 5200</li><li>• 5210</li><li>• 5230</li></ul>	<b>Vascular Solutions, Inc.</b> Minneapolis, MN	<i>There is a possibility that the excess manufacturing material may separate from the distal end of the catheter during use, and pose a potential risk of embolism.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	<b>All Unexpired Lots</b>	<b>CA</b> , nationwide	Distributed between:  <b>October 2014</b> <b>and</b> <b>September 2016</b>

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/Safety/Recalls/ucm523953.htm>